PATENT COOPERATION TREATY



PCT

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference P 26826	FOR FURTHER AC	TION See Notific Preliminary	ation of Transmittal of International Examination Report (Form PCT/IPEA/416)
International application No.	International filing date	e (day/month/year)	Priority date (day/month/year)
PCT/DE2003/003123	19 September 200	03 (19.09.2003)	19 September 2002 (19.09.2002)
International Patent Classification (IPC) or no G01N 27/30	ational classification and	I IPC	
Applicant	INFINEON TECH	NOLOGIES AG	
This international preliminary exam and is transmitted to the applicant ac This REPORT consists of a total of	ccording to Article 36.		ational Preliminary Examining Authority
This report is also accompaniamended and are the basis for 70.16 and Section 607 of the	r this report and/or sheet	ts containing rectifica	on, claims and/or drawings which have been tions made before this Authority (see Rule
These annexes consist of a to	otal of 3 s	sheets.	
3. This report contains indications rela	iting to the following iter	ms:	
I Basis of the report			
II Priority			
III Non-establishment	of opinion with regard to	o novelty, inventive st	ep and industrial applicability
IV Lack of unity of inv	vention		
V Reasoned statement citations and explar	t under Article 35(2) wit nations supporting such s	h regard to novelty, in statement	ventive step or industrial applicability;
VI Certain documents	cited		
VII Certain defects in the	he international applicati	ion	
VIII Certain observation	ns on the international ap	plication	
Date of submission of the demand		Date of completion of this report	
19 April 2004 (19.04.	.2004)	25 J	anuary 2005 (25.01.2005)
Name and mailing address of the IPEA/EP		Authorized officer	
Facsimile No.		Telephone No.	

Form PCT/IPEA/409 (cover sheet) (July 1998)

Translation

International application No.

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PCT/DE2003/003123

I. Basis	of the re	port							
1. With	regard to	the elements of the international application:*							
	the inter	mational application as originally filed							
	the desc	ription:							
K3	pages	1-27	, as originally filed						
	pages		, filed with the demand						
	pages	, filed with the letter of							
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	pages		, as originally filed						
	pages	, as amended (togethe	r with any statement under Article 19						
	pages		, filed with the demand						
ŀ	pages	1-9, filed with the letter of	30 December 2004 (30.12.2004)						
	the dra	wings: 1/4-4/4	, as originally filed						
1	pages		, filed with the demand						
ŀ	pages	, filed with the letter of							
_	pages								
╽└	the seque	ence listing part of the description:							
1	pages		, as originally filed						
	pages		, filed with the demand						
	pages	, filed with the letter of							
ماد	internationse elemen	to the language, all the elements marked above were available or furnished to to an application was filed, unless otherwise indicated under this item. In this were available or furnished to this Authority in the following language	which is:						
	the lar	nguage of a translation furnished for the purposes of international search (under I	Rule 23.1(b)).						
		nguage of publication of the international application (under Rule 48.3(b)).							
	or 55.	•							
3. Wi	ith regard	I to any nucleotide and/or amino acid sequence disclosed in the internexamination was carried out on the basis of the sequence listing:	ational application, the international						
	conta	ined in the international application in written form.							
	filed t	together with the international application in computer readable form.							
1 💄	furnis	shed subsequently to this Authority in written form.							
		shed subsequently to this Authority in computer readable form.							
	The statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.								
	_	statement that the information recorded in computer readable form is identic furnished.	al to the written sequence listing has						
4.	The a	amendments have resulted in the cancellation of:	•						
	Щ	the description, pages							
		the claims, Nos.							
		the drawings, sheets/fig							
5.	This i	report has been established as if (some of) the amendments had not been made, and the disclosure as filed, as indicated in the Supplemental Box (Rule 70.2(c)).**	since they have been considered to go						
in	this rep	nt sheets which have been furnished to the receiving Office in response to an inv ort as "originally filed" and are not annexed to this report since they do	vitation under Article 14 are referred to not contain amendments (Rule 70.16						
	nd 70.17). ny replace	ement sheet containing such amendments must be referred to under item 1 and an	nnexed to this report.						

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V.	Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability;
	citations and explanations supporting such statement

Statement			
Novelty (N)	Claims	1-9	YES
	Claims		NO
Inventive step (IS)	Claims	1-9	YES
	Claims		NO NO
Industrial applicability (IA)	Claims	1-9	YES
	Claims	· 	NO

2. Citations and explanations

This report makes reference to the following document: D1: EP-A-0 299 778.

Claim 1:

D1 discloses a method for producing a biosensor circuit
arrangement (page 5)

- in which an integrated circuit is formed in a substrate (figures 1-7),
- in which a core of an integrated reference electrode is formed by printing on the substrate with silver material as metal (example 1),
- in which subsequently the core of silver material is surrounded, at least in part, by a sleeve made of a salt of the silver material that is not readily soluble, thereby forming the integrated reference electrode (example 1),
- in which the integrated circuit is electrically coupled with the core of the integrated reference electrode (trivial feature).

Therefore, the subject matter of claim 1 differs from the known method in that

- biological molecules are applied to sensor fields of

the biosensor circuit arrangement by means of printing, whereby the sensor fields are biologically activated and whereby the printing of silver material onto the substrate and the printing of biological molecules onto the sensor fields takes place in the same operational step.

In **D1**, the reference electrode is produced in its entirety before the biological molecules are applied in a further step (see example 2).

Therefore, the subject matter of claim 1 is novel (PCT Article 33(2)).

Consequently, the problem to be solved by the present invention can be regarded as that of significantly reducing the complexity and cost of producing biosensors, since the same printing method is used in a method step in order to apply the reference electrode as that used for the biomolecules. Thereafter, the silver core is oxidized in another method step.

Therefore, the solution to this problem as proposed in claim 1 of the present application involves an inventive step (PCT Article 33(3)).

Claim 2:

Like claim 1, claim 2 contains the additional feature

that biological molecules are applied to sensor fields of the biosensor circuit arrangement by means of printing, whereby the sensor fields are biologically activated and whereby the printing of silver material onto the substrate and the printing of biological molecules onto the sensor fields takes

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place in the same operational step.

Therefore, the solution to this problem as proposed in claim 2 of the present application involves an inventive step (PCT Article 33(3)).

Dependent claims 3-9 are therefore also regarded as novel and inventive.